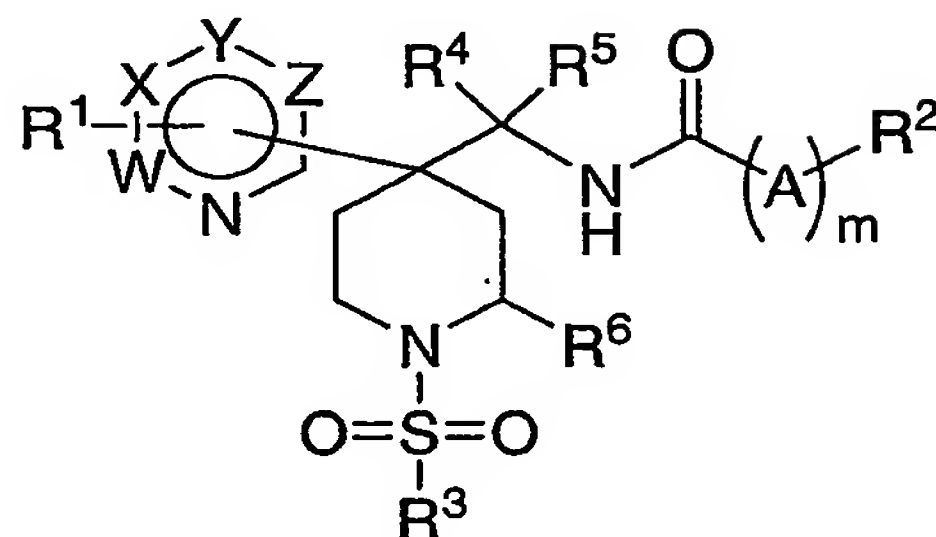


WHAT IS CLAIMED IS:

1. A compound of the formula I:



I

wherein:

R¹ is selected from one or more of the groups consisting of:

- (1) hydrogen,
- (2) C₁-6alkyl, which is unsubstituted or substituted with 1-6 halogen, hydroxy or phenyl,
- (3) -O-C₁-6alkyl,
- (4) halogen,
- (5) phenyl, which is substituted with R^{2a}, R^{2b} and R^{2c},
- (6) heterocycle, which is substituted with R^{2a}, R^{2b} and R^{2c},
- (7) -CN,
- (8) -CO₂R⁹,

wherein R⁹ is independently selected from:

- (a) hydrogen,
- (b) -C₁-6alkyl, which is unsubstituted or substituted with 1-6 fluoro,
- (c) benzyl, and
- (d) phenyl,
- (9) -SO₂R⁹,
- (10) -SO₂-NR¹⁰R¹¹,

wherein R¹⁰ and R¹¹ are independently selected from:

- (a) hydrogen,
- (b) -C₁-6alkyl, which is unsubstituted or substituted with hydroxy, 1-6 fluoro or -NR¹²R¹³, where R¹² and R¹³ are independently selected from hydrogen and -C₁-6alkyl, and where R¹⁰ and R¹¹ may be joined to form an azetidiny ring,

(c) -C₃₋₆cycloalkyl, which is unsubstituted or substituted with hydroxy, 1-6 fluoro or -NR¹²R¹³,

(d) benzyl,

(e) phenyl, and

5 (11) -CONR¹⁰R¹¹;

R² is selected from the group consisting of:

(1) phenyl, which is substituted with R^{2a}, R^{2b} and R^{2c},

(2) heterocycle, which is substituted with R^{2a}, R^{2b} and R^{2c},

10 (3) C₁₋₈alkyl, which is unsubstituted or substituted with 1-6 halogen, hydroxy, -NR¹⁰R¹¹, phenyl or heterocycle, where the phenyl or heterocycle is substituted with R^{2a}, R^{2b} and R^{2c},

(4) C₃₋₆cycloalkyl, which is unsubstituted or substituted with 1-6 halogen, hydroxy or -NR¹⁰R¹¹, and

15 (5) -C₁₋₆alkyl-(C₃₋₆cycloalkyl), which is unsubstituted or substituted with 1-6 halogen, hydroxy or -NR¹⁰R¹¹;

R^{2a}, R^{2b} and R^{2c} are independently selected from the group consisting of:

(1) hydrogen,

(2) halogen,

(3) -C₁₋₆alkyl, which is unsubstituted or substituted with:

20 (a) 1-6 halogen,

(b) phenyl,

(c) C₃₋₆cycloalkyl, or

(d) -NR¹⁰R¹¹,

(4) -O-C₁₋₆alkyl, which is unsubstituted or substituted with 1-6 halogen,

25 (5) hydroxy,

(6) -SCF₃,

(7) -SCHF₂,

(8) -SCH₃,

(9) -CO₂R⁹,

30 (10) -CN,

(11) -SO₂R⁹,

(12) -SO₂-NR¹⁰R¹¹,

(13) -NR¹⁰R¹¹,

(14) -CONR¹⁰R¹¹, and

(15) $-\text{NO}_2$;

R^3 is selected from the group consisting of:

- (1) C_{1-6} alkyl, which is unsubstituted or substituted with 1-6 halogen, hydroxyl, $-\text{NR}^{10}\text{R}^{11}$, or heterocycle, which is substituted with R^{2a} , R^{2b} and R^{2c} ,
- 5 (2) C_{3-6} cycloalkyl, which is unsubstituted or substituted with 1-6 halogen, hydroxyl or $-\text{NR}^{10}\text{R}^{11}$,
- (3) $-\text{C}_{1-6}$ alkyl- $(\text{C}_{3-6}$ cycloalkyl), which is unsubstituted or substituted with 1-6 halogen, hydroxy or $-\text{NR}^{10}\text{R}^{11}$, and
- (4) $-\text{NR}^{10}\text{R}^{11}$, and
- 10 (5) heterocycle, which is substituted with R^{2a} , R^{2b} and R^{2c} ;

R^4 and R^5 are independently selected from the group consisting of:

- (1) hydrogen, and
- (2) C_{1-6} alkyl, which is unsubstituted or substituted with halogen or hydroxyl, or R^4 and R^5 taken together form a C_{3-6} cycloalkyl ring;

15 R^6 is selected from the group consisting of:

- (1) hydrogen, and
- (2) C_{1-6} alkyl;

W, X, Y and Z are independently selected from C or N, with the proviso that at least two of W, X, Y and Z are C, to form a pyridine, oxo-dihydropyridine, pyridazine, pyrimidine, pyrazine, 1,2,4-triazine or 1,3,5-triazine ring;

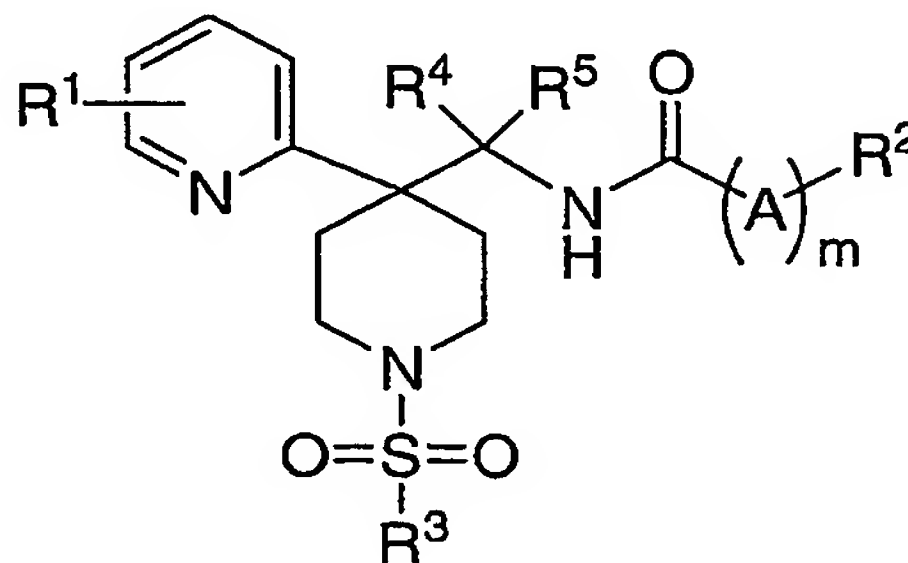
A is selected from the group consisting of:

- (1) $-\text{O}-$, and
- (2) $-\text{NR}^{10}-$;

m is zero or one, whereby when m is zero R^2 is attached directly to the carbonyl;

25 and pharmaceutically acceptable salts thereof.

2. The compound of Claim 1 of the formula Ia:



Ia

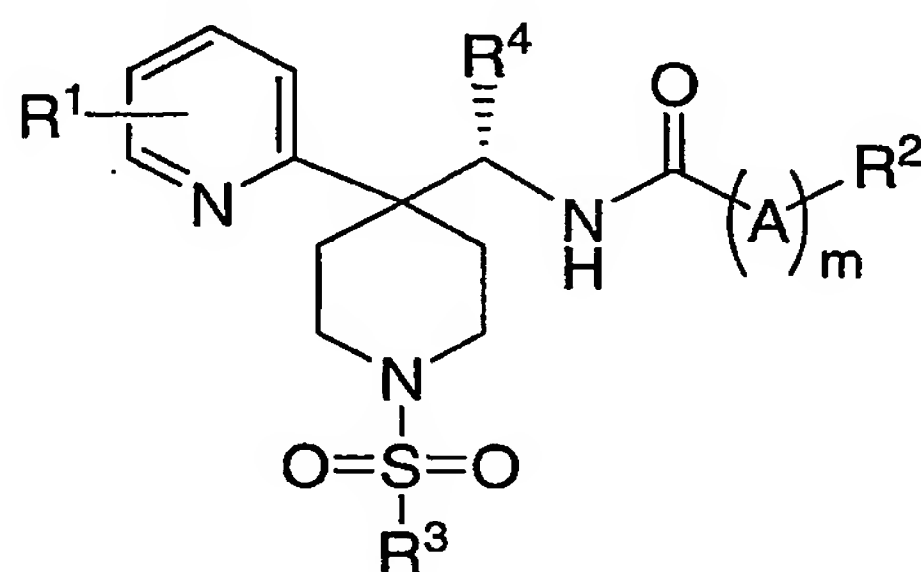
or a pharmaceutically acceptable salt thereof.

3. The compound of Claim 1 wherein R^1 is selected from the group consisting of:

- (1) hydrogen,
- (2) C_{1-3} alkyl,
- (3) fluoro,
- (4) $-CF_3$,
- (5) $-morpholinyl$, and
- (6) $-O-C_{1-3}alkyl$.

4. The compound of Claim 3 wherein R^1 is hydrogen or methyl.

5. The compound of Claim 1 of the formula Ib:



Ib

wherein R^4 is C_{1-6} alkyl; or a pharmaceutically acceptable salt thereof or an individual enantiomer or diastereomer thereof.

6. The compound of Claim 1 wherein R^4 is C_{1-3} alkyl and R^5 is hydrogen or C_{1-3} alkyl.

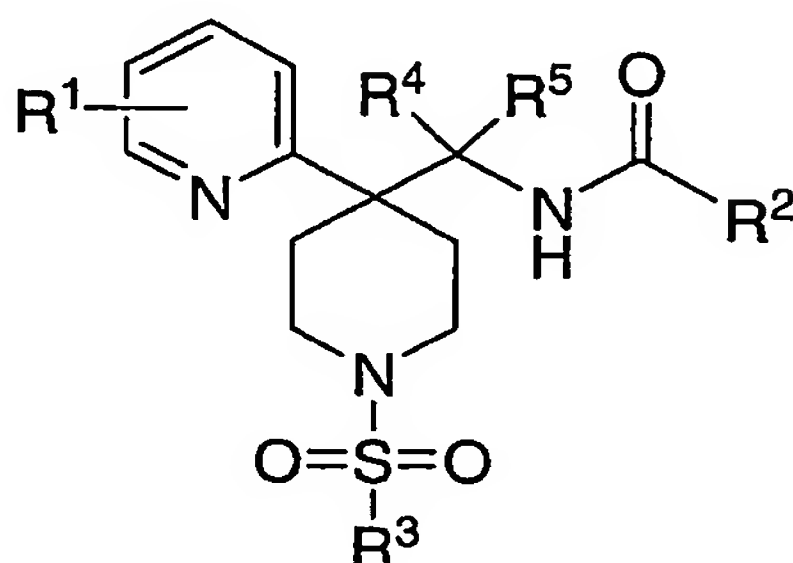
7. The compound of Claim 6 wherein R^4 is C_{1-3} alkyl in the (S) configuration and R^5 is hydrogen.

8. The compound of Claim 6 wherein R^4 is methyl and R^5 is hydrogen.

9. The compound of Claim 1 wherein R^4 is methyl and R^5 is methyl.

10. The compound of Claim 1 wherein R^4 is hydrogen and R^5 is hydrogen.

11. The compound of Claim 1 of the formula Ic:



Ic

or a pharmaceutically acceptable salt thereof.

12. The compound of Claim 1 wherein R^2 is selected from the group consisting of:

- (1) phenyl, which is substituted with R^{2a} , R^{2b} and R^{2c} ,
- (2) thienyl, which is substituted with R^{2a} , R^{2b} and R^{2c} ,
- (3) C_{1-6} alkyl, which is unsubstituted or substituted with 1-6 halogen, phenyl or $-NR^{10}R^{11}$, where the phenyl is substituted with R^{2a} , R^{2b} and R^{2c} ,
- (4) C_{3-6} cycloalkyl, which is unsubstituted or substituted with 1-6 halogen, hydroxy or $-NR^{10}R^{11}$, and

R^{2a} , R^{2b} and R^{2c} are independently selected from the group consisting of:

- (1) hydrogen,
- (2) halogen,
- (3) $-C_{1-6}$ alkyl,
- (4) $-O-C_{1-6}$ alkyl,
- (5) $-CF_3$,
- (6) $-OCF_3$,
- (7) $-OCHF_2$,
- (8) $-SCF_3$,
- (9) $-SCHF_2$, and
- (10) $-NH_2$.

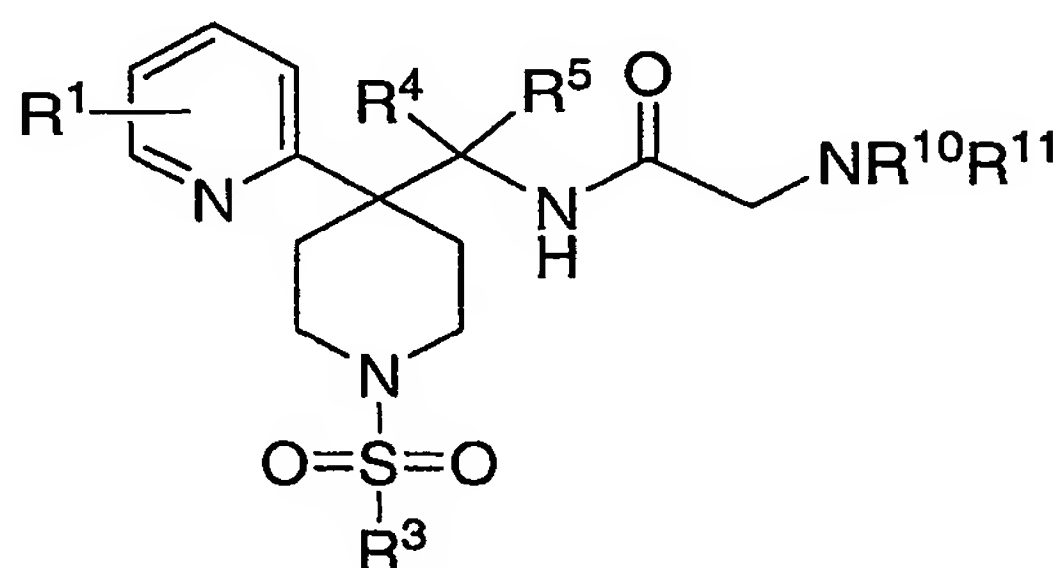
13. The compound of Claim 12 wherein R^2 is phenyl or thienyl and R^{2a} , R^{2b} and R^{2c} are independently selected from the group consisting of:

- (1) hydrogen,
- (2) halogen,
- (3) $-C_{1-6}$ alkyl,
- (4) $-O-C_{1-6}$ alkyl,
- (5) $-CF_3$,
- (6) $-OCF_3$,
- (7) $-OCHF_2$,
- (8) $-SCF_3$,
- (9) $-SCHF_2$, and
- (10) $-NH_2$.

14. The compound of Claim 13 wherein R^2 is phenyl and R^{2a} , R^{2b} and R^{2c} are independently selected from the group consisting of:

- (1) hydrogen,
- (2) fluoro,
- (3) chloro,
- (4) bromo,
- (5) $-OCH_3$,
- (6) $-CF_3$, and
- (7) $-NH_2$.

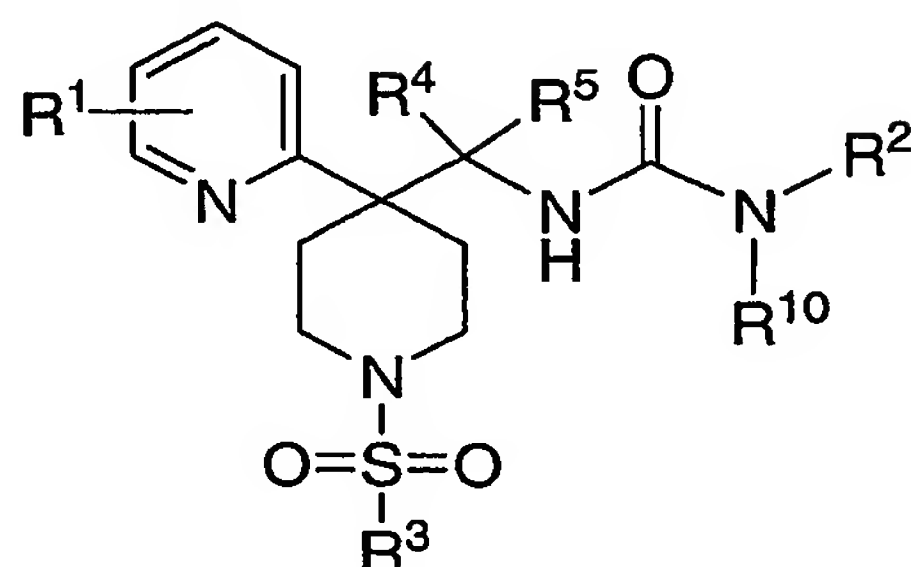
15. The compound of Claim 1 of the formula If:



If

or a pharmaceutically acceptable salt thereof.

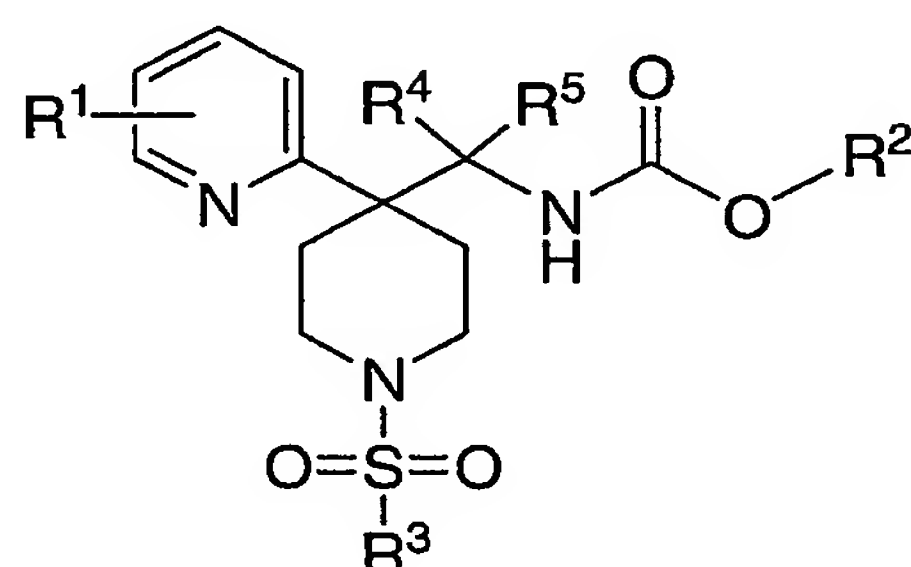
16. The compound of Claim 1 of the formula Ig:



Ig

5 or a pharmaceutically acceptable salt thereof.

17. The compound of Claim 1 of the formula Ih:



Ih

10 or a pharmaceutically acceptable salt thereof.

18. The compound of Claim 1 wherein R³ is selected from C₁-6alkyl, C₁-6alkyl-fluoro, C₃-6cycloalkyl, C₁-6alkyl-cyclopropyl, -NH(C₁-6alkyl), -N(C₁-6alkyl)(C₁-6alkyl) or azediny, which is unsubstituted or substituted with fluoro.

15

19. The compound of Claim 18 wherein R³ is -CH₂CH₃.

20. The compound of Claim 18 wherein R³ is -(CH₂)₂CH₃.

20

21. The compound of Claim 18 wherein R³ is -CH₂-cyclopropyl.

22. A compound which is selected from the group consisting of:

- 2-chloro-3,6-difluoro-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
 2,4-dichloro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
 2,4-dichloro-N-{{(1S)-1-[1-(ethylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}ethyl}benzamide;
 5 2-chloro-3,6-difluoro-N-{{1-methyl-1-[1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}ethyl}benzamide;
 2-chloro-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
 2,6-dichloro-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}-benzamide;
 2-bromo-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
 2,4-dichloro-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
 10 2-chloro-6-fluoro-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
 2-amino-6-chloro-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
 2-fluoro-6-methoxy-N-{{1-(propylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
 2-chloro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
 2-fluoro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}-6-
 15 (trifluoromethyl)benzamide;
 2,6-difluoro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
 2-chloro-6-fluoro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
 2,6-dichloro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
 2-chloro-3,6-difluoro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
 20 2-chloro-4-fluoro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
 4-chloro-2-fluoro-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide;
 2,4-dichloro-N-{{4-(4-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}benzamide
 2,4-dichloro-N-{{1-(methylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
 2,4-dichloro-N-{{1-(isopropylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide
 25 2,4-dichloro-N-{{1-(ethylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
 2,4-dichloro-N-{{1-(cyclopropylsulfonyl)-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
 2,4-dichloro-N-{{1-(propylsulfonyl)-4-pyridin-3-yl}piperidin-4-yl}methyl}benzamide;
 2,6-dichloro-N-{{1-(propylsulfonyl)-4-pyridin-3-yl}piperidin-4-yl}methyl}benzamide
 2,4-dichloro-N-{{1-(propylsulfonyl)-4-pyridin-4-yl}piperidin-4-yl}methyl}benzamide;
 30 2-chloro-6-fluoro-N-{{1-(propylsulfonyl)-4-pyridin-4-yl}piperidin-4-yl}methyl}benzamide;
 2,4-dichloro-N-{{1-[(dimethylamino)sulfonyl]-4-pyridin-2-yl}piperidin-4-yl}methyl}benzamide;
 4,4,4-trifluoro-3-methyl-N-{{4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl}methyl}butanamide;

- 2-chloro-6-fluoro-N-{[4-(6-morpholin-4-ylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl}benzamide;
2,4-dichloro-N-{[4-(6-morpholin-4-ylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl}benzamide;
2,4,5-trifluoro-N-{[4-(6-methoxypyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl}benzamide;
5 2,4-dichloro-5-fluoro-N-{[4-(6-methoxypyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl}benzamide;
N-{[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl}cyclohexanecarboxamide;
2-chloro-N-{[1-(cyclopropylsulfonyl)-4-(6-methylpyridin-2-yl)piperidin-4-yl]methyl}-3,6-difluorobenzamide;
10 N-{[1-(ethylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]methyl}-2,4-difluorobenzamide;
N-(sec-butyl)-N'-{[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]methyl}urea;
N-(4-bromophenyl)-N'-{[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]methyl}urea;
3-fluorobenzyl {[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]methyl}carbamate;
2-chlorobenzyl {[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]methyl}carbamate;
15 2,4-dichloro-N-{1-methyl-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}benzamide;
2-chloro-3,6-difluoro-N-{1-methyl-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}benzamide;
N-{1-methyl-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}-2-(trifluoromethoxy)benzamide;
2,4-dichloro-N-{1-methyl-1-[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl}benzamide;
2-chloro-3,6-difluoro-N-{1-methyl-1-[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl}benzamide;
20 2,4-dichloro-N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}benzamide;
2-chloro-3,6-difluoro-N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}benzamide;
2,4-dichloro-N-{(1S)-1-[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl}benzamide;
2-chloro-3,6-difluoro-N-{(1S)-1-[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl}benzamide;
25 4,4,4-trifluoro-3-methyl-N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}butanamide;
N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}thiophene-3-carboxamide;
N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}cyclopentane-carboxamide;
N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}cyclohexane-carboxamide;
30 2-ethyl-N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}butanamide;
2-methyl-N-{(1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl}butanamide;
3,3,3-trifluoro-N-{(1S)-1-[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl}propanamide;
2,5-dichloro-N-{(1S)-1-[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl}thiophene-3-carboxamide;

- 4-bromo-N-((1S)-1-[4-(6-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl)thiophene-3-carboxamide;
- 2,5-dichloro-N-((1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)thiophene-3-carboxamide;
- 4-bromo-N-((1S)-1-[1-(propylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)thiophene-3-carboxamide;
- 5 2-chloro-N-([1-(propylsulfonyl)-4-pyrimidin-4-ylpiperidin-4-yl]methyl)benzamide;
- 2,4-dichloro-N-([1-(propylsulfonyl)-4-pyrimidin-4-ylpiperidin-4-yl]methyl)benzamide;
- 2-chloro-3,6-difluoro-N-([1-(propylsulfonyl)-4-pyrimidin-4-ylpiperidin-4-yl]methyl)benzamide;
- 2,4-dichloro-5-fluoro-N-([1-(propylsulfonyl)-4-pyrimidin-4-ylpiperidin-4-yl]methyl)benzamide;
- 2-chloro-N-((1S)-1-[1-(ethylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)benzamide;
- 10 2-chloro-N-((1S)-1-[1-(ethylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)-3,6-difluorobenzamide;
- 2-chloro-N-((1S)-1-[1-(ethylsulfonyl)-4-(6-methylpyridin-2-yl)piperidin-4-yl]ethyl)-3,6-difluorobenzamide;
- 2,4-dichloro-N-((1S)-1-[1-(ethylsulfonyl)-4-(6-methylpyridin-2-yl)piperidin-4-yl]ethyl)benzamide;
- 2,4-dichloro-N-([1-methyl-1-[1-(azetidinesulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)benzamide;
- 15 2,4-dichloro-N-([1-methyl-1-[1-(3-fluoroazetidinesulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)benzamide;
- 2,4-dichloro-N-([1-methyl-1-[1-(3-fluoroazetidinesulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)benzamide;
- 2,4-dichloro-N-([1-methyl-1-[1-(azetidinesulfonyl)-4-(3-fluoropyridin-2-yl)piperidin-4-yl]ethyl)benzamide;
- 2,4-dichloro-N-([1-methyl-1-[1-(ethylaminosulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)benzamide;
- 20 2,4-dichloro-N-([1-[(ethylamino)sulfonyl]-4-(3-fluoropyridin-2-yl)piperidin-4-yl]methyl)benzamide;
- 2,4-dichloro-N-([1-[1-(ethylaminosulfonyl)-4-pyridin-2-ylpiperidin-4-yl]ethyl)benzamide;
- 2,4-dichloro-N-([1-[1-(ethylaminosulfonyl)-4-(3-fluoropyridin-2-yl)piperidin-4-yl]ethyl)benzamide;
- 2,4-dichloro-N-([4-(3-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl)benzamide;
- 2,4-dichloro-N-([1-(propylsulfonyl)-4-[6-(trifluoromethyl)pyridin-2-yl]piperidin-4-yl]methyl)benzamide;
- 25 2,4-dichloro-N-([1-[(cyclopropylmethyl)sulfonyl]-4-(3-methylpyridin-2-yl)piperidin-4-yl]methyl)-benzamide;
- 2,4-dichloro-N-([1-(propylsulfonyl)-4-[4-(trifluoromethyl)pyridin-2-yl]piperidin-4-yl]methyl)benzamide;
- 2,4-dichloro-N-([4-(3-chloropyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl)benzamide;
- 2,4-dichloro-N-([4-(3-methoxypyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl)benzamide;
- 30 2-chloro-N-([4-(3-chloropyridin-2-yl)-1-(ethylsulfonyl)piperidin-4-yl]methyl)-3,6-difluorobenzamide;
- 2,4-dichloro-N-([1-[(cyclopropylmethyl)sulfonyl]-4-pyridin-2-ylpiperidin-4-yl]methyl)benzamide;
- methyl [(4-[[2,4-dichlorobenzoyl]amino]methyl)-4-pyridin-2-ylpiperidin-1-yl)sulfonyl]acetate;
- 2,4-dichloro-N-([1-(ethylsulfonyl)-4-pyridin-2-ylpiperidin-4-yl]methyl)benzamide;
- 2,4-dichloro-N-([1-(propylsulfonyl)-4-pyrazin-2-ylpiperidin-4-yl]methyl)benzamide;

- 2,4-dichloro-N-(1-{1-[(3-fluoropropyl)sulfonyl]-4-pyridin-2-yl}piperidin-4-yl)ethyl)benzamide;
 2,4-dichloro-N-({1-[(3-fluoropropyl)sulfonyl]-4-pyridin-2-yl}piperidin-4-yl)methyl)benzamide;
 2,4-dichloro-N-([1-[(3-fluoropropyl)sulfonyl]-4-(3-fluoropyridin-2-yl)piperidin-4-yl]methyl)benzamide;
 2,4-dichloro-N-([1-[(cyclopropylmethyl)sulfonyl]-4-(3-fluoropyridin-2-yl)piperidin-4-yl]methyl)
 5 benzamide;
 2,4-dichloro-N-([4-(3-fluoropyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl)benzamide;
 2-chloro-3,5-difluoro-N-([1-[(cyclopropylmethyl)sulfonyl]-4-(3-fluoropyridin-2-yl)piperidin-4-yl]methyl)benzamide;
 2,4-dichloro-5-fluoro-N-([1-[(cyclopropylmethyl)sulfonyl]-4-(3-fluoropyridin-2-yl)piperidin-4-yl]methyl)benzamide;
 10 2,4-dichloro-N-([1-4-(3-fluoropyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl)benzamide;
 2,4-dichloro-N-([4-(6-trifluoromethylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl)benzamide;
 4-bromo-2-chloro-N-([1-(ethylsulfonyl)-4-(3-fluoropyridin-2-yl)-piperidin-4-yl]methyl)benzamide;
 N-(3-fluorobenzyl)-N'-([1-(propylsulfonyl)-4-(6-trifluoromethylpyridin-2-yl)piperidin-4-yl]methyl)urea;
 15 2,4-dichloro-N-([1-1-(ethylsulfonyl)-4-(3-fluoropyridin-2-yl)-piperidin-4-yl]ethyl)benzamide;
 2-bromo-4-fluoro-N-([1-(ethylsulfonyl)-4-(3-fluoropyridin-2-yl)-piperidin-4-yl]methyl)benzamide;
 2-chloro-3,6-difluoro-N-([1-(ethylsulfonyl)-4-(3-fluoropyridin-2-yl)-piperidin-4-yl]methyl)benzamide;
 2,4-dichloro-N-([1-(ethylsulfonyl)-4-(3-trifluoromethylpyridin-2-yl)-piperidin-4-yl]methyl)benzamide;
 2,4-dichloro-N-({1-[(3-fluoropropyl)sulfonyl]-4-(3-trifluoromethylpyridin-2-yl)piperidin-4-yl}methyl)benzamide;
 20 2,4,6-trifluoro-N-([1-(ethylsulfonyl)-4-(3-trifluoromethylpyridin-2-yl)-piperidin-4-yl]methyl)benzamide;
 N-(sec-butyl)-N'-([1-(propylsulfonyl)-4-(6-trifluoromethylpyridin-2-yl)piperidin-4-yl]methyl)urea;
 2,4-dichloro-N-([1-(ethylsulfonyl)-4-(3-trifluoromethylpyridin-2-yl)-piperidin-4-yl]methyl)benzamide;
 25 2,4-dichloro-N-([1-4-(3-methylpyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]ethyl)benzamide;
 N-([4-(3-bromopyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl)-2,4-dichlorobenzamide;
 2,4-dichloro-N-([4-(6-fluoropyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl)benzamide;
 2,4-dichloro-N-([4-(6-chloropyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl)benzamide;
 2,4-dichloro-N-([4-(6-oxo-1,6-dihydropyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl)benzamide;
 30 2,4-dichloro-N-([4-(3-hydroxypyridin-2-yl)-1-(propylsulfonyl)piperidin-4-yl]methyl)benzamide;
 2,4-dichloro-N-([(2[R,S],4[S,R])-2-methyl-1-(propylsulfonyl)-4-pyridin-2-yl]piperidin-4-yl]methyl)benzamide;
 2,4-dichloro-N-({1-(ethylsulfonyl)-4-[3-(trifluoromethyl)pyridin-2-yl]piperidin-4-yl}methyl)benzamide;
 or a pharmaceutically acceptable salt thereof.

23. A pharmaceutical composition which comprises a pharmaceutically acceptable carrier and a compound of Claim 1 or a pharmaceutically acceptable salt thereof.

5 24. A method for inhibiting the glycine transporter GlyT1 in a mammal in need thereof which comprises the administration of an effective amount of the compound of Claim 1 or a pharmaceutically acceptable salt thereof.

10 25. A method for the manufacture of a medicament for inhibiting the glycine transporter GlyT1 in a mammal in need thereof comprising combining the compound of Claim 1 or a pharmaceutically acceptable salt thereof with a pharmaceutical carrier or diluent.

15 26. A method for treating a neurological and psychiatric disorders associated with glycinergic or glutamatergic neurotransmission dysfunction in a mammalian patient in need thereof which comprises administering to the patient a therapeutically effective amount of a compound of Claim 1 or a pharmaceutically acceptable salt thereof.

20 27. A method for treating schizophrenia in a mammalian patient in need thereof which comprises administering to the patient a therapeutically effective amount of a compound of Claim 1 or a pharmaceutically acceptable salt thereof.